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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-13-0612]

Proposed Data Collections Submitted for
Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404-639-7570 and send comments to Ronald Otten, CDC Reports Clearance Officer, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an email to omb@cdc.gov.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the

collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Written comments should be received within 60 days of this notice.

Proposed Project

Well-Integrated Screening and Evaluation for Women Across the Nation (WISEWOMAN) Reporting System (OMB #0920-0612, exp. 3/31/2013) - Extension - National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Cardiovascular disease (CVD), which includes heart disease, myocardial infarction, and stroke, is the leading cause of death for women in the United States, and is largely preventable. The WISEWOMAN program (Well-Integrated Screening and Evaluation for Women Across the Nation), administered by the Centers for Disease Control and Prevention (CDC), was established to examine ways to improve the delivery of services for women who have limited access to health care and elevated risk factors for CVD. The program focuses on reducing CVD risk factors and provides screening services for select risk factors such as elevated blood cholesterol, hypertension and abnormal blood glucose levels. The

program also provides lifestyle interventions and medical referrals. On an annual basis, 21 grantees funded through the WISEWOMAN program have provided services to approximately 30,000 women who are already participating in the National Breast and Cervical Cancer Early Detection Program (NBCCEDP), also administered by CDC.

CDC currently collects information from WISEWOMAN grantees to support continuous program monitoring and improvement activities. CDC seeks to extend OMB approval for one additional year. There are no changes to the number of respondents, the data items reported to CDC, the estimated burden per response, or the total estimated annualized burden. All information will continue to be collected twice per year.

Information reported to CDC includes baseline and follow-up data (12 months post enrollment) for all women served through the WISEWOMAN program. These data, called the minimum data elements (MDE), include data elements that describe risk factors for the women served in each program and data elements that describe the number and type of intervention sessions attended. Funded grantees compile the data from their existing databases and report the MDE to CDC on April 15th and October 15th of each year. The MDE data provide an assessment of how effective the WISEWOMAN

program is at reducing the burden of cardiovascular disease risk factors among women who utilize program services. The information collected from grantees is also used to assess the cost-effectiveness and impact of the program. Because certain demographic information has already been collected as part of NBCCEDP, the additional burden of WISEWOMAN program reporting is modest.

The overall program evaluation is designed to demonstrate how WISEWOMAN can obtain more complete health data on vulnerable populations, promote public education about disease incidence and risk-factors, improve the availability of screening and diagnostic services for under-served women, ensure the quality of services provided to under-served women, and develop strategies for improved interventions. The information reported to CDC also includes programmatic information related to grantee management, public education and outreach, professional education, service delivery, cost, and progress toward meeting stated programmatic objectives.

All MDE information will be submitted to CDC electronically.

The estimated burden per response for Screening and Assessment

MDE is 16 hours. The estimated burden per response for Lifestyle

Intervention MDE is 8 hours. Progress reports will be submitted

in hardcopy format. The estimated burden per response for each progress report is 16 hours.

Estimated Annualized Burden Hours

Type of Respondent	Form Name	Number of Respondents	Number of Responses per Respondent	Average Burden per Response (in hr)	Total Burden (in hr)
WISEWOMAN Grantees	Screening and Assessment MDE	21	2	16	672
	Lifestyle Intervention MDE	21	2	8	336
	Progress Report	21	2	16	672
				Total	1,680

DATE: October 9, 2012

Ron A. Otten,

Director, Office of Scientific Integrity (OSI)

Office of the Associate Director for Science (OADS)

Office of the Director

Centers for Disease Control and Prevention

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